L Number	Hits	Search Text	DB	Time stamp
- Livallibei	1524	angiogenic ADJ factor	USPAT;	Time stamp 2003/01/08 10:41
	1324	angiogenic ADJ factor		2003/01/08 10:41
			US-PGPUB;	
			EPO; JPO;	
	23010	chimeric	DERWENT	2002/01/00 10:41
-	23010	Chimieric	USPAT;	2003/01/08 10:41
			US-PGPUB;	
			EPO; JPO;	
	145333	ahima aria arr 6 mia r	DERWENT	2002/04/02 40 44
-	145232	chimeric or fusion	USPAT;	2003/01/08 10:41
			US-PGPUB;	
			EPO; JPO;	
		(aminomia ADI forton) anno ablass (DERWENT	2000/04/00 40 40
-	9	(angiogenic ADJ factor) same chimeric	USPAT;	2003/01/08 10:46
			US-PGPUB;	
			EPO; JPO;	
	22012		DERWENT	
-	23013	vegf adj fusion or chimeric	USPAT;	2003/01/08 10:47
			US-PGPUB;	
			EPO; JPO;	
	, <u>-</u>		DERWENT	
-	17	vegf adj (fusion or chimeric)	USPAT;	2003/01/08 10:48
			US-PGPUB;	
			EPO; JPO;	
			DERWENT	
-	. 10	fgf adj (fusion or chimeric)	USPAT;	2003/01/08 10:48
			US-PGPUB;	
			EPO; JPO;	
			DERWENT	
-	46	vegf\$ adj (fusion or chimeric)	USPAT;	2003/01/08 10:59
			US-PGPUB;	
			EPO; JPO;	
			DERWENT	
-	0	vaegf-A and (vegf\$ adj (fusion or chimeric))	USPAT;	2003/01/08 10:59
			US-PGPUB;	
			EPO; JPO;	
			DERWENT	
-	0	vaegf-A121 and (vegf\$ adj (fusion or chimeric))	USPAT;	2003/01/08 11:00
1			US-PGPUB;	
			EPO; JPO;	
			DERWENT	
-	0	vaegf-A or VEGF-A121 OR VEGF-A145 OR	USPAT;	2003/01/08 11:01
	į	VEGF-A165 OR VEGF-A189 OR VEGF-A206	US-PGPUB;	·
	İ		EPO; JPO;	
	l		DERWENT	
-	174	vegf-A or VEGF-A121 OR VEGF-A145 OR	USPAT;	2003/01/08 11:03
		VEGF-A165 OR VEGF-A189 OR VEGF-A206	US-PGPUB;	
			EPO; JPO;	
	İ		DERWENT	
-	152	(vegf-A or VEGF-A121 OR VEGF-A145 OR	USPAT;	2003/01/08 11:02
	1	VEGF-A165 OR VEGF-A189 OR VEGF-A206) AND	US-PGPUB;	
	ļ	(chimeric or fusion)	EPO; JPO;	
			DERWENT	
-	0	(vegf-A or VEGF-A121 OR VEGF-A145 OR	USPAT;	2003/01/08 11:02
		VEGF-A165 OR VEGF-A189 OR VEGF-A206) NEAR	US-PGPUB;	
	ļ	(chimeric or fusion)	EPO; JPO;	
			DERWENT	
-	9	(vegf-A or VEGF-A121 OR VEGF-A145 OR	USPAT;	2003/01/08 11:03
		VEGF-A165 OR VEGF-A189 OR VEGF-A206) SAME	US-PGPUB;	
		(chimeric or fusion)	EPO; JPO;	
		•	DERWENT	

-	218	vegf-B or VEGF-B167 OR VEGF-B186	USPAT;	2003/01/08 11:22
		_	US-PGPUB;	
			EPO; JPO;	
			DERWENT	
-	205	vegf-d	USPAT;	2003/01/08 11:03
		7 - 5 - 5	US-PGPUB;	2005,01,00 11.05
			EPO; JPO;	
			DERWENT	
_	134	vegf-E	USPAT;	2003/01/08 11:03
-	134	vegrat	•	2003/01/08 11.03
			US-PGPUB;	
			EPO; JPO;	
	1	(ACIDIC ADI FIRDORI ACT. ADI CDOUTI ADI	DERWENT	
-	1551	(ACIDIC ADJ FIBROBLAST ADJ GROWTH ADJ	USPAT;	2003/01/08 11:04
		FACTOR) OR AFgf	US-PGPUB;	
			EPO; JPO;	
			DERWENT	
-	2918	(BASIC ADJ FIBROBLAST ADJ GROWTH ADJ	USPAT;	2003/01/08 11:05
		FACTOR) ORBFgf	US-PGPUB;	
			EPO; JPO;	
			DERWENT	
-	540007	ANGIOPOIETIN-1 OF ang1	USPAT;	2003/01/08 11:05
		<u>-</u>	US-PGPUB;	
	[EPO; JPO;	•
			DERWENT	
-	13	vegf-d SAME (chimeric or fusion)	USPAT;	2003/01/08 11:05
		(3)	US-PGPUB;	
			EPO; JPO;	
			DERWENT	
_	5	vegf-E SAME (chimeric or fusion)	USPAT;	2003/01/08 11:06
		(children of fasion)	US-PGPUB;	2003/01/00 11:00
			EPO; JPO;	
			DERWENT	
_	50	((ACIDIC AD) FIBROBLAST ADJ GROWTH ADJ	USPAT;	2003/01/08 11:06
	30	FACTOR) OR AFgf) SAME (chimeric or fusion)	US-PGPUB;	2003/01/08 11.00
		TACTOR) OR ALGIT SAME (CHIMENE OF TUSION)		
			EPO; JPO; DERWENT	
_	70	(/BASIC AD1 EIBDOBLAST AD1 CDOWTH AD1	ſ	2002/01/09 11:06
_	/ /	((BASIC ADJ FIBROBLAST ADJ GROWTH ADJ	USPAT;	2003/01/08 11:06
		FACTOR) ORBFgf) SAME (chimeric or fusion)	US-PGPUB;	
			EPO; JPO;	
	10	(ANCIODOIETIN 1 OF prod) CAME (-blos)	DERWENT	2002/01/02 11 22
-	13	(ANGIOPOIETIN-1 OF ang1) SAME (chimeric or	USPAT;	2003/01/08 11:06
		fusion)	US-PGPUB;	
			EPO; JPO;	
			DERWENT	
-	12	(vegf-B or VEGF-B167 OR VEGF-B186) SAME	USPAT;	2003/01/08 11:07
		(chimeric or fusion)	US-PGPUB;	
			EPO; JPO;	
			DERWENT	
-	6	VEGF-B167	USPAT;	2003/01/08 11:23
			US-PGPUB;	
			EPO; JPO;	
			DERWENT	

were significantly less necrotic, suggesting that necrosis in these tumors is the result of insufficient angiogenesis. ANSWER 2 OF 3 CAPLUS CO-RIGHT 2003 ACS L9 ACCESSION NUMBER: DOCUMENT NUMBER: 128:71142 Targeting the tumor vasculature: inhibition of tumor TITLE: growth by a vascular endothelial growth factor-toxin AUTHOR(S): Olson, Timothy A.; Mohanraj, D.; Roy, Sabita; Ramakrishnan, S. Department of Pharmacology, University of Minnesota, **CORPORATE SOURCE:** Minneapolis, MN, USA International Journal of Cancer (1997), 73(6), 865-870 CODEN: IJCNAW; ISSN: 0020-7136 SOURCE: **PUBLISHER:** Wiley-Liss, Inc. DOCUMENT TYPE: Journal LANGUAGE: English Tumor-derived vascular endothelial growth factor (VEGF)/vascular permeability factor (VPF) plays an important role in neovascularization and the development of tumor stroma. Furthermore, VEGF receptors are over-expressed in the endothelial cells of tumor vasculature and almost non-detectable in the vascular endothelium of adjoining normal tissues. The differential expression of receptor offers a selective advantage for targeting cytotoxic toxin polypeptides. We have prepd. a vascular targeting reagent by chem. linking recombinant VEGF to a truncated form of diphtheria toxin. The VEGF-toxin conjugate was selectively toxic to endothelial cell lines and inhibited exptl. neovascularization of the chick chorioallantoic membrane. In the present study, we examd. the effects of VEGF-toxin conjugate on solid tumor growth. Athymic nude mice with established s.c. tumors were treated with daily i.p. injections of the VEGF-toxin conjugate or free toxin. When compared with control animals treated with the toxin polypeptide alone, the conjugate-treated animals displayed a significant inhibition of tumor growth. Histol. anal. of tumors from conjugate-treated animals revealed hemorrhagic necrosis consistent with a vascular-mediated injury. In contrast, highly vascularized normal tissues from conjugate-treated animals demonstrated no evidence of hemorrhage or tissue injury. The conjugate was well tolerated without apparent toxicities. Our results illustrate the anti-tumor activity of a VEGF-toxin conjugate selectively targeting the tumor neovasculature. L9 ANSWER 3 OF 3 COPYRIGHT 2003 Univentio PCTFULL **** DATA NOT AVAILABLE FOR THIS ACCESSION NUMBER **** DATA NOT AVAILABLE FOR THIS ACCESSION NUMBER => d his (FILE 'HOME' ENTERED AT 16:59:28 ON 08 JAN 2003) FILE 'MEDLINE' ENTERED AT 16:59:40 ON 08 JAN 2003 L1 6036 S VEGF? L2 102848 S FUSION L3 59 S L1 (S) L2 220976 S TARGET? L4 L5 126030 S ENDOTHELI? 14 S L1 (S) L2 (S) L4 (S) L5 FILE 'BIOSIS, EMBASE, SCISEARCH, CAPLUS, PCTFULL' ENTERED AT 17:05:43 ON 08 JAN 2003 **L7**

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59 DUP REM L7 (27 DUPLICATES REMOVED)
              3 S L8 NOT PY>1998
=> logoff hold
COST IN U.S. DOLLARS
                                                   SINCE FILE
                                                                   TOTAL
                                                        ENTRY
                                                                 SESSION
FULL ESTIMATED COST
                                                        17.86
                                                                   24.81
DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)
                                                   SINCE FILE
                                                                   TOTAL
                                                        ENTRY
                                                                 SESSION
CA SUBSCRIBER PRICE
                                                        -0.65
                                                                   -0.65
SESSION WILL BE HELD FOR 60 MINUTES
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STN INTERNATIONAL SESSION SUSPENDED AT 17:09:24 ON 08 JAN 2003

86 S L6

L8

transgeniques exprimant ces proteines et des analogues fonctionnellement equivalents de ces proteines. L'invention cerne enfin des methodes permetant d'induire la differenciation de motoneurones somatiques et de traiter des maladies liees a la carence en motoneurones fonctionnant normalement, des maladies neurodegeneratives, des troubles neurologiques et des maladies neuromusculaires.

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ANSWER 5 OF 5
L17
                                                                                                    DUPLICATE 2
                                         MEDLINE
                                    95403738
ACCESSION NUMBER:
                                                           MEDLINE
DOCUMENT NUMBER:
                                    95403738
                                                        PubMed ID: 7673487
TITLE:
                                    Transplanted human neurons derived from a teratocarcinoma
                                    cell line (NTera-2) mature, integrate, and survive for over
                                    1 year in the nude mouse brain.
                                    Kleppner S R; Robinson K A; Trojanowski J Q; Lee V M
AUTHOR:
                                   Department of Pathology and Laboratory Medicine, University of Pennsylvania Medical School, Philadelphia 19104, USA.
CORPORATE SOURCE:
                                    JOURNAL OF COMPARATIVE NEUROLOGY, (1995 Jul 10) 357 (4)
SOURCE:
                                    618-32.
                                    Journal code: 0406041. ISSN: 0021-9967.
PUB. COUNTRY:
                                    United States
                                    Journal; Article; (JOURNAL ARTICLE)
DOCUMENT TYPE:
LANGUAGE:
                                   English
FILE SEGMENT:
                                   Priority Journals
ENTRY MONTH:
                                   199510
ENTRY DATE:
                                   Entered STN: 19951026
                                   Last Updated on STN: 19970203
                                   Entered Medline: 19951019
        Retinoic acid (RA) induces a human teratocarcinoma cell line (NTera-2 or
AB
        NT2) to give rise exclusively to ***post*** - ***mitotic***

***neuron*** -like (NT2N) cells, but NT2N cells never acquire a fully mature neuronal phenotype in vitro. To determine whether NT2N cells can mature into adult neuron-like cells in vivo, purified NT2N cells were grafted into different regions of the central nervous system (CNS) of
        adult and neonatal athymic mice, and the grafts were examined immunohistochemically by light, confocal, and electron microscopy using
         antibodies to a panel of developmentally regulated neuronal polypeptides.
        NT2N grafts were distinguished from endogenous mouse neurons with
        antibodies that recognize human or murine specific epitopes in selected neuronal polypeptides. Viable NT2N cells were identified in > 89% of graft recipients (N = 90), and some grafts survived 14 months. Within 3 weeks of ***implantation***, grafted NT2N cells re-extended their processes, and
        the location of the grafts (e.g., septum versus neocortex) appeared to
        determine the extent to which processes were elaborated. Within the early
         post-transplantation period, grafted NT2N cells expressed the same
        neuronal polypeptides as their in vitro counterparts. However, between 6
        weeks and 4-6 months post- ***implantation***, the grafted NT2N cells progressively acquired the molecular phenotype of fully mature in vivo neurons as evidenced by dramatically increased expression of the most highly phosphorylated isoforms of the heavy neurofilament subunit, and the de novo expression of adult CNS tau. Notably, the time course for the extension of processes and the expression of neuronal polypeptides by NT2N grafts was similar in peopatal and adult mice. Although grafted NT2N cells
        grafts was similar in neonatal and adult mice. Although grafted NT2N cells formed synapse-like structures and elaborated dendrites and axons, these
        axons remained unmyelinated. Finally, none of the transplanted NT2N cells
        reverted to a neoplastic state. These studies demonstrate that pure populations of grafted human NT2N cells acquire a fully mature neuronal phenotype in vivo, and that these cells integrate and survive for > 1 year post- ***implantation*** in the mouse CNS. These human neuron-like cells are an attractive model system for studies of neuronal development,
        polarity and transplantation.
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(FILE 'HOME' ENTERED AT 09:52:39 ON 08 JAN 2003)
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L8
          68237 S ALZHEIMERS DISEASE OR PARKINSONS DISEASE OR HUNTINGTONS DISEA
L9
         123638 S ALZHEIMER? DISEASE OR PARKINSON? DISEASE OR HUNTINGTON? DISEA
L10
        -229826 S POST-MITOTIC
                                  URON OR NEURON
         344762 S REVIEW
            171 S L3 (S) L6 (S) L9
13 S L3 (S) L6 (S) L9 (S) L11
L13
L14
              0 S L3 (S) L6 (S) L9 (S) L11 (S) L10
              8 S L3 (S) L6 (S) L11 (S) L10
L15
     FILE 'PCTFULL, USPATFULL, MEDLINE, BIOSIS, EMBASE, CAPLUS, CONFSCI,
     SCISEARCH' ENTERED AT 10:17:36 ON 08 JAN 2003
L16
              9 S L5
L17
              5 DUP REM L16 (4 DUPLICATES REMOVED)
=> logoff hold
COST IN U.S. DOLLARS
                                                   SINCE FILE
                                                                    TOTAL
                                                        ENTRY
                                                                  SESSION
FULL ESTIMATED COST
                                                        18.50
                                                                    39.12
 SESSION WILL BE HELD FOR 60 MINUTES
STN INTERNATIONAL SESSION SUSPENDED AT 10:20:47 ON 08 JAN 2003
Connecting via Winsock to STN
Welcome to STN International! Enter x:X
LOGINID:ssspta1632rrs
PASSWORD:
TERMINAL (ENTER 1, 2, 3, OR ?):2
* * * * * * * * * *
                       Welcome to STN International
NEWS
                  Web Page URLs for STN Seminar Schedule - N. America
                   "Ask CAS" for self-help around the clock
 NEWS
          Apr 08
          Apr 09
 NEWS
                  BEILSTEIN: Reload and Implementation of a New Subject Area
          Apr 09
 NEWS
                  ZDB will be removed from STN
          Apr
 NEWS
              19
                  US Patent Applications available in IFICDB, IFIPAT, and IFIUDB
 NEWS
          Apr
                  Records from IP.com available in CAPLUS, HCAPLUS, and ZCAPLUS
          Apr 22
 NEWS
                  BIOSIS Gene Names now available in TOXCENTER
          Apr 22
                  Federal Research in Progress (FEDRIP) now available
 NEWS
          Jun 03
 NEWS
                  New e-mail delivery for search results now available
 NEWS 10
          Jun 10
                  MEDLINE Reload
 NEWS 11
          Jun 10
                  PCTFULL has been reloaded
 NEWS 12
          Jul 02
                  FOREGE no longer contains STANDARDS file segment
          Jul 22
 NEWS 13
                  USAN to be reloaded July 28, 2002;
                  saved answer sets no longer valid
 NEWS 14
          Jul 29
                  Enhanced polymer searching in REGISTRY
                  NETFIRST to be removed from STN CANCERLIT reload
          Jul 30
 NEWS
      15
          Aug 08
 NEWS 16
          Aug 08
 NEWS 17
                  PHARMAMarketLetter(PHARMAML) - new on STN
NEWS 18
          Aug 08
                  NTIS has been reloaded and enhanced
 NEWS 19
          Aug 19
                  Aquatic Toxicity Information Retrieval (AQUIRE)
                  now available on STN
          Aug 19
NEWS 20
                  IFIPAT, IFICDB, and IFIUDB have been reloaded
NEWS 21
          Aug 19
                  The MEDLINE file segment of TOXCENTER has been reloaded
 NEWS
      22
          Aug 26
                  Sequence searching in REGISTRY enhanced
      23
 NEWS
          Sep 03
                  JAPIO has been reloaded and enhanced
NEWS 24
                  Experimental properties added to the REGISTRY file
          Sep 16
NEWS 25
          Sep 16
                  Indexing added to some pre-1967 records in CA/CAPLUS
          Sep 16
                  CA Section Thesaurus available in CAPLUS and CA
 NEWS 26
NEWS 27
          Oct 01
                  CASREACT Enriched with Reactions from 1907 to 1985
 NEWS 28
          Oct 21
                  EVENTLINE has been reloaded
 NEWS 29
          Oct 24
                  BEILSTEIN adds new search fields
NEWS 30
          Oct 24
                  Nutraceuticals International (NUTRACEUT) now available on STN
NEWS
          Oct 25
                  MEDLINE SDI run of October 8, 2002
 NEWS
          Nov 18
                  DKILIT has been renamed APOLLIT
          Nov 25
NEWS 33
                  More calculated properties added to REGISTRY
                  TIBKAT will be removed from STN
NEWS 34
          Dec 02
          Dec 04
                  CSA files on STN
NEWS 35
NEWS 36
          Dec 17
                  PCTFULL now covers WP/PCT Applications from 1978 to date
NEWS 37
          Dec 17
                  TOXCENTER enhanced with additional content
NEWS 38
          Dec 17
                  Adis Clinical Trials Insight now available on STN
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